

Bryan G Hughes

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

816
citations

759233

12
h-index

996975

15
g-index

21
all docs

21
docs citations

21
times ranked

1531
citing authors

#	ARTICLE	IF	CITATIONS
1	Evolutionary conservation of the clk-1-dependent mechanism of longevity: loss of mclk1 increases cellular fitness and lifespan in mice. <i>Genes and Development</i> , 2005, 19, 2424-2434.	5.9	309
2	Sequential fractionation and isolation of subcellular proteins from tissue or cultured cells. <i>MethodsX</i> , 2015, 2, 440-445.	1.6	145
3	Targeting MMP-2 to treat ischemic heart injury. <i>Basic Research in Cardiology</i> , 2014, 109, 424.	5.9	69
4	A Mild Impairment of Mitochondrial Electron Transport Has Sex-Specific Effects on Lifespan and Aging in Mice. <i>PLoS ONE</i> , 2011, 6, e26116.	2.5	45
5	MMP-2 is localized to the mitochondria-associated membrane of the heart. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2014, 306, H764-H770.	3.2	40
6	Different Mechanisms of Longevity in Long-Lived Mouse and <i>Caenorhabditis elegans</i> Mutants Revealed by Statistical Analysis of Mortality Rates. <i>Genetics</i> , 2016, 204, 905-920.	2.9	37
7	Genetic and molecular characterization of CLK-1/mCLK1, a conserved determinant of the rate of aging. <i>Experimental Gerontology</i> , 2006, 41, 940-951.	2.8	33
8	Estimating the occurrence of primary ubiquinone deficiency by analysis of large-scale sequencing data. <i>Scientific Reports</i> , 2017, 7, 17744.	3.3	31
9	Nuclear matrix metalloproteinase-2 in the cardiomyocyte and the ischemic-reperfused heart. <i>Journal of Molecular and Cellular Cardiology</i> , 2016, 94, 153-161.	1.9	30
10	Many possible maximum lifespan trajectories. <i>Nature</i> , 2017, 546, E8-E9.	27.8	25
11	Dynamic Alterations to $\hat{\alpha}$ -Actinin Accompanying Sarcomere Disassembly and Reassembly during Cardiomyocyte Mitosis. <i>PLoS ONE</i> , 2015, 10, e0129176.	2.5	21
12	Matrix metalloproteinase-2 in oncostatin M-induced sarcomere degeneration in cardiomyocytes. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2016, 311, H183-H189.	3.2	16
13	Doxorubicin induces de novo expression of N-terminal-truncated matrix metalloproteinase-2 in cardiac myocytes. <i>Canadian Journal of Physiology and Pharmacology</i> , 2018, 96, 1238-1245.	1.4	12
14	Compensatory elevation of voluntary activity in mouse mutants with impaired mitochondrial energy metabolism. <i>Physiological Reports</i> , 2014, 2, e12214.	1.7	2
15	Mclk1 ^{+/-} mice are not resistant to the development of atherosclerosis. <i>Lipids in Health and Disease</i> , 2009, 8, 16.	3.0	1
16	Phylogenetic ubiquity of the effects of altered ubiquinone biosynthesis on survival. <i>Aging</i> , 2011, 3, 184-185.	3.1	0
17	Role of MMP-2 activation in oncostatin M induced cardiomyocyte dedifferentiation. <i>FASEB Journal</i> , 2013, 27, 1146.4.	0.5	0
18	Analysis of mitochondrial MMP-2 and MMP-9 in the heart. <i>FASEB Journal</i> , 2013, 27, 1129.10.	0.5	0

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19	Intracellular proteases and sarcomere disassembly in neonatal cardiomyocytes. FASEB Journal, 2013, 27, 1217.33.	0.5	0
20	Matrix metalloproteinaseâ€2 is localized to the mitochondriaâ€associated membrane in the heart (1154.4). FASEB Journal, 2014, 28, 1154.4.	0.5	0
21	Nuclear Localization and Biological Function of Matrix Metalloproteinaseâ€2. FASEB Journal, 2015, 29, 979.6.	0.5	0